Application No. 10/536,533
Paper Dated: August 24, 2009
In Reply to USPTO Correspondence of May 22, 2009
Attorney Docket No. 4544-051675

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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

Claims 1-22 (Cancelled)

Claim 23 (Currently Amended): A process for preparing an agglutination reagent for detecting typhoid comprising:

- (a) preparing a polyclonal-monospecific antibody specific to Salmonella typhi;
- (b) preparing a latex particle suspension; and
- (c) coating a latex particle with said polyclonal-monospecific antibody specific to Salmonella typhi;

wherein said polyclonal-monospecific antibody specific to Salmonella typhi is prepared according to a method comprising:

- raising a hyper immune sera against a purified protein encoded by a Flagellin gene specific to Salmonella typhi, and
- (ii) separating said polyclonal-monospecific antibody fraction—from said hyper immune sera:

wherein said latex particle suspension is prepared according to a method comprising:

- (i) mixing 1% carboxylated latex particles of—size—and a 40 mM 2-N morpholinoethane sulphonic acid (MES) buffer of pH 5.5 to 6.0 in a ratio of 1:1, washing with a 20 mM MES buffer of pH 5.5 thereby forming a washed latex particle, and
- (ii) adding a 1-ethyl-3 (3-dimethyl-amino propyl) carbodiimide hydrochloride (EDC) in a 20 mM MES buffer of pH 5.5 to said washed latex particle in a ratio of 1:1, washing with a 20 mM MES buffer (pH 5.5); and

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- (a) preparing a polyclonal-monospecific antibody specific to Salmonella typhi;
- (b) preparing a latex particle suspension; and
- (c) coating a latex particle with said polyclonal-monospecific antibody specific to Salmonella typhi;

wherein said polyclonal-monospecific antibody specific to Salmonella typhi is prepared according to a method comprising:

- raising a hyper immune sera against a purified protein encoded by a Flagellin gene specific to Salmonella typhi, and
- separating said polyclonal-monospecific antibody fraction—from said hyper immune sera;

wherein said latex particle suspension is prepared according to a method comprising:

- (i) mixing 1% carboxylated latex particles of—size—and a 40 mM 2-N morpholinoethane sulphonic acid (MES) buffer of pH 5.5 to 6.0 in a ratio of 1:1, washing with a 20 mM MES buffer of pH 5.5 thereby forming a washed latex particle, and
- (ii) adding a 1-ethyl-3 (3-dimethyl-amino propyl) carbodiimide hydrochloride (EDC) in a 20 mM MES buffer of pH 5.5 to said washed latex particle in a ratio of 1:1, washing with a 20 mM MES buffer (pH 5.5); and